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ning of each regular issue of the PCT Gazette.

(54) Title: METHODS FOR THE IDENTIFICATION OF AGENTS THAT MODULATE THE STRUCTURE AND PROCESSING
OF A MEMBRANE BOUND PRECURSOR PROTEIN

(57) Abstract: The present invention provides methods for the screening and identification of agents from a large library of molecu-
lar structures that can alter the cleavage of a membrane protein of interest. Agents identified by the methods of the present invention
that modify the cleavage of the membrane protein can be used in the treatment and prevention of diseases such as inflammation,
diabetes, cancer, Alzheimer's disease, Parkinson's disease, and the like. The methods select for and identify effector agents that
bind to the membrane protein of interest causing a structural change in the structure of the membrane protein in such a way that the
efficiency of the cleavage of a secretase is modulated. Further, the methods are carried out in an in vivo system that provides for
physiological conditions similar or identical to conditions for membrane protein processing. Agents can be selected for their ability
to cause a decrease or increase the amount of secretase cleavage of the membrane protein.

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INTERNATIONAL SEARCH REPORT

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A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C12N 15/00; G01N 33/53, 33/567; A61K 49/00
 US CL : 435/7.1, 7.21, 172.3; 530/350; 800/3

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

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 U.S. : 435/7.1, 7.21, 172.3; 530/350; 800/3

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
 EAST, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X --- Y	US 6,440,698 B1 (GURNEY et al) 27 August 2002 (27.08.2002), column 10, lines 30-59; column 7, lines 34-46; column 35, line 7 to column 36, line 21; Table 4.	1-7, 36-38, 46 ----- 15-35, 39-45, 47-54
X --- Y	HOOPER, N.M. et al. Membrane Protein Secretases. Biochem. J. 1997, Vol 321. pages 265-279, especially pages 268-272 (sections on (-amyloid precursor protein secretases and ACE secretase), and 275-276 (section on Secretase Assays, Whole Cell Assays).	1-7, 36-38 ----- 15-35, 39-45, 47-54
X --- Y	US 6,175,057 B1 (MUCKE et al) 16 January 2001 (16.01.2001), column 16, lines 1-58; column 17, lines 1-16.	1, 8-14 ----- 42-45, 53-54
Y	US 2002/0127564 A1 (NOLAN) 12 September 2002 (12.09.2002), entire document, especially page 2, paragraph 25 and 27; page 3, paragraphs 37-38; page 4, paragraphs 41-45; page 5, paragraphs 51-53; page 11, paragraphs 105-107.	15-20, 22-35, 47-54



Further documents are listed in the continuation of Box C.



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INTERNATIONAL SEARCH REPORT

C. (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 6,420,110 B1 (GYURIS et al) 16 July 2002 (16.07.02) entire document, especially column 2, lines 31-33; column 19, lines 22-29; column 43, lines 22-24; column 18, lines 11-29; column 9, lines 36-37.	21, 24, 28, 32, 45, 52
Y	RIAN, E. et al. A Signal Sequence Trap Based on Cell Enrichment Using Anti-CD19 Antibody Coated Magnetic Beads. Scand J Immunol. September 2001. Vol 54, pages 280-284, entire document.	26
Y	PONCET, C. CD24, a glycoposphatidylinositol-anchored molecule is transiently expressed during the development of human central nervous system and is a marker of human neural cell lineage tumors. Acta Neuropathol. 1996, Vol. 91, No. 4, Abstract only.	30
Y	MURTHY, S.C. et al. Characterization of the interleukin 3 receptor. Exp Hematol. January 1990, Vol. 18, No. 1, Abstract only.	31
Y	MAZUR-KOLECKA, B. et al. Accumulation of Alzheimer amyloid-peptide in cultured myocytes is enhanced by serum and reduced by cerebrospinal fluid. J Neuropathol Exp Neurol. March 1997, Vol. 56, No. 3, Abstract only.	39-41, 51, 54